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<ul> <li>To combine searches use # before search number, e.g., #2 AND #6.</li> <li>Search numbers may not be continuous; all searches are represented.</li> </ul>			
Entrez		<b>:</b>	
PubMed	Search Most Recent Queries	Time	ole committee from the matter control of the contro
	#49 Search Swanson S and detection Limits: Publication Date to 1999/08/25	14:33:18	<u>23</u>
	#48 Search Swanson S and adenovirus Limits: Publication Date to 1999/08/25	14:32:39	<u>1</u>
PubMed Services	#47 Search Swanson S Limits: Publication Date to 1999/08/25	14:32:29	<u>326</u>
	#46 Search Mytych D Limits: Publication Date to 1999/08/25	14:32:08	<u>. 2</u>
	#45 Search peptide linker and CKGKG Limits: Publication Date to 1999/08/25	14:26:12	1195
	#44 Search peptide linker Limits: Publication Date to 1999/08/25	14:25:57	<u>1195</u>
	#43 Search CKGKG Field: All Fields, Limits: Publication Date to 1999/08/25	14:25:36	0
Related Resources	#36 Search Lemon S and HCV Limits: Publication Date to 2000/12/23	09:59:45	<u>22</u>
	#35 Search Lemon S Limits: Publication Date to 2000/12/23	09:46:17	<u>174</u>
	#34 Search HCV replicon and tat Limits: Publication Date to 2000/12/23	09:43:39	<u>0</u>
	#30 Search transactivation and HCV Field: All Fields, Limits: Publication Date to 2000/12/23	09:40:14	13
	#28 Search Guo 2001 and HCV	09:14:26	<u>5</u>
	#27 Search Guo 2000 and HCV	09:14:08	<u>4</u>
	#25 Search Blight 2000 and hcv	09:08:04	<u>2</u>
	#23 Search Bartenschlager 2000 and hcv	09:00:41	<u>4</u>
	#20 Search Lohmann 1999 and HCV	08:46:00	<u>6</u>
	#18 Search Lohmann 2001 and HCV	08:42:26	<u>7</u>
	#16 Search Pietschamnn 2001 and self-replication Field: All Fields, Limits: Publication Date to 2000/12/23	08:40:04	<u>12</u>
	#15 Search Pietschamnn 2001 and self-replication	08:39:23	<u>27</u>
	#14 Search Pietschamnn 2001 and HCV RNA	08:39:12	<u>1059</u>
	#13 Search Pietschamnn 2001 and HCV	08:38:37	<u>2577</u>
	#11 Search Reynolds 1995 and HCV	08:37:24	<u>2</u>
	#9 Search Reynolds 1996 and HCV	08:36:45	<u>1</u>

```
=> "peptide linker"
        305983 "PEPTIDE"
        223298 "PEPTIDES"
        391308 "PEPTIDE"
                  ("PEPTIDE" OR "PEPTIDES")
         14823 "LINKER"
          3477 "LINKERS"
         16810 "LINKER"
                  ("LINKER" OR "LINKERS")
           359 "PEPTIDE LINKER"
L5
                  ("PEPTIDE"(W) "LINKER")
=> antigen and L1
        249705 ANTIGEN
        198079 ANTIGENS
        309817 ANTIGEN
                  (ANTIGEN OR ANTIGENS)
L6
            87 ANTIGEN AND L1
=> assay and L6
        300651 ASSAY
        128138 ASSAYS
        392648 ASSAY
                  (ASSAY OR ASSAYS)
L7
             4 ASSAY AND L6
=> adenovirus and L6
         19990 ADENOVIRUS
          2838 ADENOVIRUSES
         20494 ADENOVIRUS
                  (ADENOVIRUS OR ADENOVIRUSES)
^{\text{L8}}
              0 ADENOVIRUS AND L6
```

REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 3 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1997:238402 CAPLUS

DOCUMENT NUMBER: 126:220707

TITLE: Multimer compositions for conferring immunogenicity to

a peptide

Stanton, G. John; Hughes, Thomas K., Jr.; Smith, Eric INVENTOR (S):

Μ.

PATENT ASSIGNEE(S): Board of Regents, the University of Texas System, USA

SOURCE: PCT Int. Appl., 115 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE -----\_\_\_\_ -----WO 9705886 A1 19970220 WO 1996-US12632 19960805 W: AU, CA, JP RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE US 5807552 A 19980915 US 1995-511662 19950804 AU 9666452 A1 19970305 AU 1996-66452 19960805 A1 PRIORITY APPLN. INFO.: US 1995-511662 19950804

WO 1996-US12632

19960805

## ABSTRACT:

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A multimer of monomers non-covalently held together by interactive \*\*\*peptide\*\*\* linkers if provided for the enhancement of the immunogenicity of a substance. These multimers are useful for stimulating or suppressing the immune system, detecting the presence of antibodies, bypassing MHC restriction in an animal, and the effective presentation of antigen , suppressing autoimmune disease, inducing cytokine production, adsorption, treating a defective immune system and for use as an adjuvant. The invention specifically describes multimers in which monomers are peptide sequences containing an HIV HP-6 epitope with left- and right-flanking linker sequences.

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1996:162202 CAPLUS

DOCUMENT NUMBER: 124:229436

TITLE: Production and characterization of bispecific

single-chain antibody fragments

De Jonge, Jan; Brissinck, Jan; Heirman, Carlo; Demanet, Christian; Leo, Oberdan; Moser, Muriel; AUTHOR (S):

Thielemans, Kris

Lab. Physiol., Med. Sch. Vrije Univ. Brussel, CORPORATE SOURCE:

Brussels, B-1090, Belg.

Molecular Immunology (1995), 32(17/18), 1405-12 CODEN: MOIMD5; ISSN: 0161-5890 SOURCE:

PUBLISHER: Elsevier Journal DOCUMENT TYPE: English LANGUAGE:

ABSTRACT:

We report the construction, expression and purification of a bispecific single-chain Fv antibody fragment produced in Escherichia coli. The protein possesses a dual specificity: the single-chain FvB1 portion is directed to the Idiotype of BCL1 lymphoma cells, the single-chain Fv2C11 moiety binds to the CD3 marker on T cells. The two domains are joined by a flexible peptide Using Immobilized Metal Affinity Chromatog., the recombinant \*\*\*linker.\*\*\* protein was purified from bacterial insol. membrane fractions. After refolding of the bispecific protein, it was affinity-purified. As demonstrated by flow

cytometry, both binding sites are retained in the refolded protein. Retargeted cytotoxicity and T cell proliferation **assays** further prove the biol. activity and specificity of the bispecific single-chain Fv. Thus, these bispecific mols. show a potential antitumor activity.

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